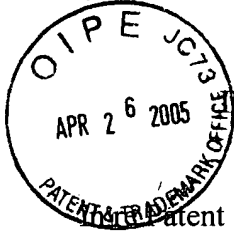


02047.000154.

  
PATENT



IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Patent of:

INDRA PRAKASH

Appl. No.: 09/859,439

Filed: May 18, 2001

For: SYNTHESIS OF N-[N-(3,3-DIMETHYLBUTYL)-L- $\alpha$ -ASPARTYL]-L-PHENYLALANINE  
1-METHYL ESTER USING  
OXAZOLIDINONE DERIVATIVES

U.S. Patent No.: US 6,852,875 B2

Issued: February 8, 2005

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Examiner: Paul A. Zucker

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Group Art Unit: 1621

)  
:  
April 25, 2005

Commissioner for Patents  
P.O. Box 1450  
Alexandria, VA 22313-1450

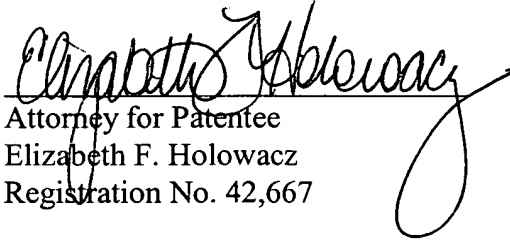
CERTIFICATE OF CORRECTION UNDER RULE 322

Sir:

It is respectfully requested that a Certificate of Correction be issued by the Patent and Trademark Office due to errors which appear in the printed patent as a result of Patent and Trademark Office mistakes. A Certificate of Correction form, in duplicate, is enclosed.

Patentee's undersigned attorney may be reached in our New York office by telephone at (212) 218-2100. All correspondence should continue to be directed to our address given below.

Respectfully submitted,



Attorney for Patentee  
Elizabeth F. Holowacz  
Registration No. 42,667

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UNITED STATES PATENT AND TRADEMARK OFFICE  
CERTIFICATE OF CORRECTION

PATENT NO. : US 6,852,875 B2

DATED : February 8, 2005

INVENTOR(S) : INDRA PRAKASH

Page 1 of 1

It is certified that error appears in the above-identified patent and that said Letters Patent is hereby corrected as shown below:

ON COVER PAGE (56) REFERENCES CITED

Other Publications, "Chemmiker Zeitlung 1990," should read --Chemiker Zeitung 1990,--.

COLUMN 7:

Line 25, "acid." should read --acid. ¶

Neo-aspartic acid (5 mmol) was dissolved in 2,2-dimethoxypropane (10 ml) and 1,4-dioxane (10 ml). p-Toluenesulfonic acid (0.5 mmol) was added to the reaction mixture and refluxed for 48 hours. The solvent was removed from the reaction mixture, extraction using dichloromethane was performed, the organic layer was concentrated by vacuo and the residue was checked via <sup>1</sup>H NMR. 2-[(4S)-3-(3,3-dimethylbutyl)-2,2-dimethyl-5-oxo-1, 3-oxazolan-4-yl]acetic acid was obtained in about a 20% yield and with low purity.--.

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